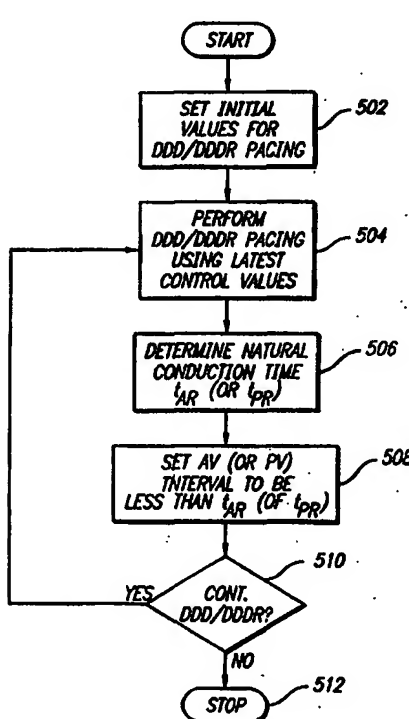


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(21) International Application Number: PCT/US98/17506 (22) International Filing Date: 24 August 1998 (24.08.98) (30) Priority Data: 08/919,430 27 August 1997 (27.08.97) US (71) Applicant: PACESETTER, INC. [US/US]; 15900 Valley View Court, Sylmar, CA 91392-9221 (US). (72) Inventors: LEVINE, Paul, A.; 23452 Windcrest Place, Newhall, CA 91321 (US). VALIKAI, Kenneth; 27226 Sunny Ridge Road, Palos Verdes Pen., CA 90274 (US). (74) Agent: ROMANO, Malcolm, J.; Pacesetter, Inc., 15900 Valley View Court, Sylmar, CA 91392-9221 (US).		(81) Designated States: European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>With international search report.</i>
(54) Title: DUAL CHAMBER IMPLANTABLE PACEMAKER HAVING NEGATIVE AV/PV HYSTERESIS AND ECTOPIC DISCRIMINATION (57) Abstract <p>A DDD-mode or DDDR-mode pacer operating with set initial parameters (502, 504) measures the natural conduction time of the patient (506), and then automatically sets the AV or PV interval to be less than that measured time (508) to avoid operating the pacer in response to an ectopic heartbeat.</p>  <pre>graph TD START([START]) --> 502[SET INITIAL VALUES FOR DDD/DDDR PACING] 502 --> 504[PERFORM DDD/DDDR PACING USING LATEST CONTROL VALUES] 504 --> 506[DETERMINE NATURAL CONDUCTION TIME (AR (OR) PR)] 506 --> 508[SET AV (OR) PV INTERVAL TO BE LESS THAN (AR (OR) PR)] 508 --> 510{CONT. DDD/DDDR?} 510 -- YES --> 504 510 -- NO --> 512([STOP])</pre>		

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**DUAL-CHAMBER IMPLANTABLE PACEMAKER HAVING
NEGATIVE AV/PV HYSTERESIS AND ECTOPIC DISCRIMINATION**

Field of the Invention

The present invention relates to implantable
5 medical devices and methods, and more particularly to
an implantable pacemaker that automatically adapts its
atrial-ventricular (AV) delay to a value related to
the measured natural conduction time of a patient,
where the measured natural conduction time is measured
10 in a way that ignores ectopic R-waves. Such pacemaker
is particularly suited for maximizing the cardiac
output of patients suffering from hypertrophic
obstructive cardiomyopathy.

Background of the Invention

15 The heart is a pump that pumps life-sustaining
blood throughout the body of the patient. A good
summary of how the heart performs its function of a
pump, including a brief description of the physiology
of the heart, may be found in U.S. Patent 5,340,361,
20 incorporated herein by reference.

A pacemaker is an implantable medical device that
monitors the activity of the heart for the occurrence
of P-waves and/or R-waves, and steps in with
electronically generated stimuli, when needed, to
25 force the depolarization of the atria and/or
ventricles. A pacemaker-generated stimulus that is
delivered to the atrium is referred to herein as an
"A-pulse". A pacemaker-generated stimulus that is
delivered to the ventricle is referred to herein as a
30 "V-pulse". Most pacemakers are configured to provide
an A-pulse and/or V-pulse only if a prescribed period
of time has elapsed without the occurrence of a P-wave

and/or an R-wave, i.e., without the occurrence of natural heartbeats.

The prescribed period of time used by the pacemaker between contraction of the ventricle and contraction of the atrium is generally referred to as the V-A Interval or the atrial escape interval. For most dual-chamber pacemaker modes of operation, only if a P-wave does not occur during the atrial escape interval will the pacemaker step in at the conclusion of such interval and generate an A-pulse.

The prescribed period of time used by the pacemaker between contraction of the atrium and contraction of the ventricle is referred to as the A-V Interval, or sometimes it is called the "AV Delay". The pacemaker, for most dual-chamber modes of operation, generates a V-pulse only if the AV Delay elapses without the occurrence of an R-wave.

In the above-described manner, the heart is thus afforded as much time as possible to beat on its own before the electronically-generated stimuli of the pacemaker are delivered to the heart, causing it to beat at the rate set by the pacemaker.

Heretofore, most cardiac patients using a pacemaker have suffered from at least one of various cardiac conditions or diseases that affect either the ability of the SA node to maintain and sustain a satisfactory heartbeat rate (hereafter "rate problems"), or the ability of the AV node or the AV bundle to conduct a suitable stimulus to the ventricle (hereafter "conduction problems"). Advantageously, both rate problems and conduction problems lend themselves well to a pacemaker solution because the underlying cardiac muscle tissue is in place and is capable of responding to the electronically-generated stimuli produced by the pacemaker.

In recent years, it has also been recognized that standard dual-chamber pacemakers can also be effective

in treating individuals who have a primary abnormality of cardiac muscle function with totally normal rhythms. While there are subsets of cardiomyopathy, each of which has its own specific therapeutic regimens, dual-chamber pacing with a short AV delay intentionally resulting in a disordered sequence of ventricular activation, has been very effective in alleviating symptoms of chest pain, shortness of breath and syncope in patients with hypertrophic obstructive cardiomyopathy (HOCM) that has proven to be refractory to pharmacologic therapy.

In HOCM, there is a disproportionate thickening of the intraventricular septum resulting in a dynamic outflow tract obstruction with hypercontractility of the remaining muscle. This obstruction to the ejection of blood from the ventricle causes the heart muscle to work harder increasing its metabolic demand resulting in episodes of chest pain identical to angina pectoris but without concomitant atherosclerotic obstruction of the coronary arteries, shortness of breath or dyspnea and recurrent syncope. The primary therapy for this disorder is pharmacologic; using drugs which intentionally decrease the vigor of contraction. The net effect is that the degree of obstruction is reduced, improving the overall cardiac efficiency. Until recently, the only available option when pharmacologic therapy was no longer effective was an open heart operation to surgically excise a portion of the abnormally thickened muscular septum. As this is a procedure performed infrequently by most cardiac surgeons, the operative morbidity and even mortality was higher than for most cardiac operations. In recent years, it has been recognized that a disordered sequence of ventricular activation (as might occur with the spontaneous development of an intraventricular conduction abnormality called Left Bundle Branch Block

(LBBB)) can delay the electrical activation of the intraventricular septum, referred to simply as the septum, thus delaying its mechanical contraction and reducing the outflow tract obstruction. The net
5 effect is a reduction in the major symptoms associated with HOCM. In that a similar ventricular activation sequence can be induced by pacing from the right ventricular apex, dual-chamber pacing has been
10 successfully employed to also treat these individuals who do not have any conduction system problem that would otherwise constitute a standard indication for pacing.

In accordance with the present invention, a dual-chamber pacemaker is implanted in patients suffering
15 from HOCM and is programmed with a sufficiently short AV or PV interval to preempt the normal conduction pattern and initiate the desired disordered sequence of ventricular activation from the location of the ventricular pacing lead. During PV or AV pacing, the
20 pacemaker delivers a V-pulse to the ventricles at a programmed delay after the occurrence of the atrial event, which atrial event could be either the occurrence of a P wave or the delivery of an A-pulse. With delivery of a V-pulse to the tissue before
25 natural conduction is allowed to take place, the pacemaker causes electrical depolarization and resultant muscular contraction pattern that is different from the normal pattern. This altered mechanical contraction sequence, for many HOCM
30 patients, advantageously results in a reduction in the degree of outflow tract gradient, improving cardiac output and reducing the work of the heart. The result is a reduction in the severity and incidence of the symptoms of chest pain, dyspnea and syncope.
35 PV or AV pacing is only effective, however, when the V-pulse is delivered to the ventricular tissue before the occurrence of an R-wave, i.e., before the

ventricular tissue depolarizes. As soon as the ventricular tissue depolarizes, it becomes refractory, and will not respond to a V-pulse, until such time as it repolarizes. It is thus necessary, if AV or PV
5 pacing is to be used, to set the AV (or PV) interval of the pacemaker to a value that is less than the patient's normal conduction time. Heretofore, this requirement has forced the AV (or PV) interval to be set to very short values, i.e., between 80 and 120 ms
10 or shorter, because during exercise (or other periods of physical activity or physiological stress) the patient's native conduction time may shorten significantly. Thus, in order to guarantee that the pacemaker will always pace the ventricles (i.e., in
15 order to assure that the V-pulse is delivered to the ventricular tissue at a time when it is not refractory), the AV (or PV) interval must be set to an interval that is shorter than any native conduction interval that might exist in the given patient at any
20 given time.

Disadvantageously, setting a very short programmed AV (or PV) interval may adversely affect cardiac output because it may force ventricular contraction well before the ventricles have had
25 sufficient time to be filled with blood from the atrium. Thus, what is needed for patients suffering from a HOCM is a pacemaker that paces the ventricles at a time in the cardiac cycle that is always less than the natural conduction time, i.e., at a time that
30 is prior to the occurrence of an R-wave, but that is not so much less than the natural conduction. That is, what is needed is a pacemaker that automatically sets its internally-generated AV and/or PV intervals to be just short of the patient's native conduction
35 time, thereby assuring that the AV (or PV) interval is sufficiently long to allow the blood to physically move from the atrium to the ventricles; yet remains

sufficiently short to always be less than the patient's native conduction time, thereby assuring that the V-pulse is not delivered when the ventricular tissue is refractory.

5 Existing systems, see, e.g., U.S. Patent No. 5,340,361, incorporated herein by reference, respond to a single sensed R-wave by shortening the subsequent AV or PV interval by a programmable interval or delta. However, an ectopic R-wave that fails to represent
10 true conduction, but rather a coincidental event following a paced or sensed atrial event, will result in a shortening of the AV or PV interval for the ensuing 256 cycles. The shortened interval may be physiologically inappropriate and cause hemodynamic
15 deterioration.

 When the AV and/or PV interval is shortened to a value that is just less than the native conduction time, there is also a need to regularly check the interval to determine whether it is still set at a
20 value that is just less than the patient's native conduction time. This is because the patient's native conduction time will normally change over a period of time and may increase without detection, leaving the conduction time too short. A particular stair-step
25 searching technique for regularly checking the AV and/or PV interval to determine if it is still less than the native conduction time is described in U.S. Patent No. 5,334,220, which patent is incorporated herein by reference. It would also be desirable to
30 employ other techniques or systems wherein the AV or PV interval is always maintained at an interval that bears the prescribed relationship to the patient's natural conduction time, thereby obviating the need to periodically "search" for such an optimum value.

35 Some additional techniques known in the art for carrying out ventricular pacing, and in particular, for verifying the complete ventricular capture results

from application of a V-pulse, are shown in U.S. Patent Nos. 5,507,782 and 5,514,163.

Summary of the Invention

The present invention addresses the above and
5 other needs by providing a dual-chamber programmable
pacemaker, and method of operating such a pacemaker,
that adaptively adjusts the pacemaker's AV interval to
assure that the AV interval remains a fixed amount
less than a measured natural conduction time while
10 ignoring ectopic R-waves. The pacemaker is used to
implement a method of treating a patient suffering
from hypertrophic obstructive cardiomyopathy (HOCM)
through preemptive ventricular pacing.

In a first embodiment of the invention, the
15 pacemaker includes an atrial sense amplifier,
ventricular sense amplifier, a pulse generator, and a
control system. As is common with conventional dual-
chamber pacemakers, the atrial sense amplifier senses
P-waves in an atrial channel. The P-waves represent
20 natural atrial activity. The ventricular sense
amplifier senses R-waves in the ventricular channel.
The R-waves represent natural ventricular activity.
The pulse generator generates an atrial stimulation
pulse (A-pulse) in the atrial channel in the absence
25 of a sensed P-wave by the atrial sense amplifier
within an atrial escape interval. The pulse generator
also generates a ventricular stimulation pulse (V-
pulse) in the ventricular channel in the absence of a
sensed R-wave by the ventricular sense amplifier
30 within an AV/PV interval. The control system defines
the AV/PV interval and the atrial escape interval.
The AV/PV interval begins upon the sensing of a P-wave
or the generation of an A-pulse, whichever event
occurs first in the atrial channel. The atrial escape
35 interval begins upon the sensing of an R-wave or the

generation of a V-pulse, whichever event occurs first in the ventricular channel.

Unlike conventional pacemakers, however the pacemaker provides for ventricular pacing by setting the AV/PV time interval to a value that is less than a measured conduction time interval by a prescribed amount, and takes extra precautions to assure that the measured conduction time is not tainted by the occurrence of an ectopic R-wave. A timer, which is part of the control system, periodically measures the conduction time interval using techniques that minimize the influence of ectopic R-waves on such conduction time measurement. The conduction time interval is the time period between atrial activity in the atrial channel and the sensing of an R-wave in the ventricular channel. The atrial activity includes the generation of an A-pulse or the sensing of a P-wave in the atrial channel, whichever event occurs first in a given cardiac cycle.

In a first aspect of the invention, the control system **decreases** the AV/PV interval by a first prescribed interval (e.g., by 20 ms) after a first prescribed number of consecutive cardiac cycles in which a sensed R-wave occurs within the AV time interval. In one embodiment, the first prescribed number of consecutive cardiac cycles in which the control system looks to find an R-wave in the AV-PV interval is preferably at least 3 cardiac cycles, however, this could be a programmable value selected by the physician. In this way, a single ectopic R-wave will not trigger a decrease to an unphysiologic AV interval and the hemodynamics of the patient will be preserved.

Secondly, the occurrence of an R-wave resulting from natural A-V conduction may be verified by monitoring the conduction time interval (e.g., for at

least three, or a programmable number of, consecutive sensed R-waves) and computing an average of the conduction time interval measurements thus made. Because it is highly unlikely that three or even two
5 consecutive ectopic R-waves would ever occur, the natural conduction time measurement is thus not likely to be tainted by ectopic R-waves if the measurement is based on the occurrence of, e.g., at least three consecutive R-waves of similar timing. The control
10 system, after the detection of a verified R-wave, decreases the AV/PV time interval by the first prescribed interval such that the AV/PV interval of the pacemaker is set to a value that is from 5-100 ms less than the average conduction time interval.

15 Further, the control system may decrease the AV/PV time interval only if the natural conduction time for each R-wave of the last three consecutive sensed R-waves exceeds a predetermined conduction time range of plus or minus 25 ms of the average conduction
20 time. Again, this helps ensure that an ectopic R-wave (which is not likely to meet this criteria) will not be considered.

In a second aspect of the invention, the control system **increases** the AV/PV time interval by a second
25 prescribed interval (e.g., by 20 ms) after a second prescribed number of consecutive cardiac cycles without a sensed R-wave. This process is repeated until an R-wave occurs. Thereafter, when the first prescribed number of consecutive cardiac cycles occurs
30 **with** a sensed R-wave falling within the AV time interval, then the AV/PV time interval is again decreased to a value that is less than the measured conduction time interval by the first prescribed interval.

35 In order to sort out ectopic R-waves, so that such ectopic R-waves do not adversely influence or

taint the conduction time measurement, a more detailed feature of the present invention requires that the second prescribed number of consecutive cardiac cycles which must occur **without** a sensed R-wave before the control system increases the second prescribed interval comprises either a preset value (e.g., at least 3 cardiac cycles) or programmable value (e.g., between 4 and 512 cardiac cycles, corresponding to approximately 4.0 seconds and 8.5 minutes if pacing at 60 bpm). An isolated ectopic R-wave, if one occurs, would break this specified string of consecutive cardiac cycles without an R-wave. However, as repetitive, consecutive, sensed R-waves are required to define the conduction interval before the AV interval is shortened, isolated ectopic R-waves (even occurring coincidentally within the AV time interval) will not adversely affect the behavior of the pacing system.

In a second embodiment of the invention, the programmable pacemaker includes a timing circuit, a programmable processor, and sensing circuitry. The timing circuit defines a conduction time interval between a paced or sensed atrial event and a natural ventricular contraction. The programmable processor sets and derives operating parameters of the pacemaker, including an AV/PV interval defined by the timing circuit. As mentioned, the AV/PV interval is the time interval between a sensed P-wave representing natural atrial contractions or generation of an A-pulse, representing an atrial stimulation pulse, and a V-pulse representing a ventricular stimulation pulse, in absence of a sensed R-wave. The sensing circuitry electrically senses atrial and ventricular depolarizations. The processor reduces an initial value of the AV/PV interval by a predetermined time interval in response to at least three consecutive

sensed R-waves, indicative of repetitive natural ventricular contractions, and returns the AV/PV interval to its initial value after a predetermined number of ventricular depolarizations.

5 In another similar embodiment of the present invention, the pacemaker similarly includes a sensing circuitry, a timing circuit, and a programmable processor. The sensing circuitry electrically senses P-waves and R-waves that are associated with natural
10 atrial and ventricular contractions, respectively. The timing circuit measures a conduction time interval between a P-wave or a sensed atrial event and an R-wave or a paced ventricular event. The programmable processor sets and derives operating parameters of the
15 pacemaker, including the AV/PV interval. The AV/PV interval is set to a value that is a prescribed time interval less than the last measured conduction time interval. Upon the sensing of an R-wave, the processor compares the sensed R-wave with a sensed R-
20 wave stored in the processor's memory to verify that the sensed R-wave is the result of a conducted ventricular contraction and is not the result of an ectopic ventricular contraction. The processor decreases the value of the AV/PV interval by a first
25 predetermined time interval in response to a predetermined number of consecutive verified R-waves sensed by the sensing circuitry, indicative of natural ventricular contractions.

30 In other more detailed features of the invention, the processor may select the reference

 R-wave from a plurality of R-wave morphologies stored in the processor. More particularly, the processor may select the reference R-wave under external control through a telemetry link.

35 Alternatively, the pacemaker may further include a physiological sensor that generates a signal representing the patient's physiological condition,

and the processor may automatically select the reference R-wave based on the patient's state signal.

Brief Description of the Drawings

The above and other aspects, features and
5 advantages of the present invention will be more apparent from the following more particular description thereof, presented in conjunction with the following drawings wherein:

FIG. 1 is a timing diagram indicating preemptive
10 ventricular pacing for treatment of hypertrophic obstructive (HOCM) cardiomyopathy exhibiting immediate AV/PV interval adjustment in response to a single R-wave;

FIG. 2A is a timing diagram indicating preemptive
15 ventricular pacing for treatment of HOCM exhibiting AV/PV interval adjustment in response to three consecutive R-waves, in accordance with the present invention;

FIG. 2B is a timing diagram indicating preemptive
20 ventricular pacing for treatment of HOCM cardiomyopathy exhibiting no AV/PV interval adjustment in response to a single R-wave, in accordance with the present invention;

FIG. 2C is a timing diagram indicating preemptive
25 ventricular pacing for treatment of HOCM exhibiting no AV/PV interval adjustment in response to two consecutive R-waves, in accordance with the present invention;

FIG. 3A is a graph showing a representative shape
30 or morphology of an R-wave resulting from natural AV conduction.

FIG. 3B is a graph showing various representative measured natural and ectopic R-waves for morphology comparison with the standard R-wave of **FIG. 3A**;

FIG. 4 is a schematic cross-sectional representation of a healthy heart;

FIG. 5 is a schematic cross-sectional representation of a heart suffering from HOCM;

5 FIG. 6 is a functional block diagram of a representative multi-mode microprocessor-based implantable pacemaker that may be used with the present invention;

FIG. 7 is a flowchart illustrating a method including the present invention; and

FIG. 8 is a more detailed flowchart illustrating the method of the present invention when used to provide preemptive ventricular pacing to treat HOCM.

Corresponding reference characters indicate corresponding components throughout the several views of the drawings.

Detailed Description of the Invention

The following description is of the best mode presently contemplated for carrying out the invention.

20 This description is not to be taken in a limiting sense, but is made merely for the purpose of describing the general principles of the invention. The scope of the invention should be determined with reference to the claims.

25 As indicated above, the present invention is directed to an implantable dual-chamber pacemaker, and a method of operating an implantable dual-chamber pacemaker, that maintains its AV interval (or PV interval) at a value that is less than the heart's
30 natural conduction time and that thereafter automatically adapts or adjusts the AV/PV interval based on the detection of, or lack of detection of, R-waves.

More specifically, if the pacemaker detects and verifies the occurrence of R-waves, indicating a shortening of the natural conduction time interval, the pacemaker shortens the AV/PV interval by a prescribed time interval. After a prescribed number of cardiac cycles without detecting an R-wave, the pacemaker lengthens the AV/PV interval by a prescribed time interval, until the AV/PV interval is at its initial value or until the next periodic measurement of the natural conduction time.

The natural conduction time is measured by the pacemaker as the time interval between an atrial event, which could either be a P-wave or an A-pulse, and an R-wave. Throughout the remaining description of the invention, unless the context indicates otherwise, the natural conduction time (whether measured from a P-wave or an A-pulse) is generically referred to as t_{AR} . Once the natural conduction time has been determined, the AV (or PV) interval of the pacemaker is set to a value that is a slightly less than t_{AR} , e.g., t_{AR} minus Δ , where Δ is a programmable number of from, e.g., 5-100 ms. The natural conduction time is thereafter periodically determined, e.g., every x cardiac cycles (where x is a prescribed number, e.g., from 10 to 1024) by the sensing circuits of the pacemaker, so that the AV (or PV) interval can be regularly updated to remain just less than the most-recently determined t_{AR} .

The pacemaker reduces the potentially deleterious effects of ectopic R-waves by several methods. One method involves requiring the occurrence of at least three consecutive A-R or P-R complexes (hereinafter simply referred to as an "R-wave") before reducing the AV/PV interval. This method is further described with reference to **FIGS. 1 through 2C** which show timing diagrams of cardiac cycles. Each cardiac cycle is

shown commencing with a P-wave 52, representing natural atrial activity. Each P-wave is initially followed by a V-pulse 53, representing a ventricular stimulation pulse. If V-pulse 53 is of sufficient energy, it captures the ventricle and is followed by a paced R-wave 54. For purposes of the present invention, and the discussion that follows, it is assured that a V-pulse 53 always captures the ventricle, thereby causing a paced R-wave 54 to occur immediately after the V-pulse. For the sake of clarity, a paced R-wave 54 is shown as a negative-going pulse while a naturally occurring (non-paced)

R-wave 56, representing natural ventricular activity, is represented by a positive-going pulse.

FIG. 1 depicts the operation of a conventional pacemaker having an adaptive AV/PV interval. See, e.g., U.S. Patent No. 5,340,361. Such pacemaker shortens the AV/PV interval in response to only one R-wave 56. More particularly, as can be determined in **FIG. 1**, the AV/PV interval is initially set, based on the natural conduction time, to 250 ms and upon the occurrence of a single

R-wave 56 after a conduction time interval t_{AR} of only 200 ms, the AV/PV interval is shortened to 170 ms.

However, as shown in **FIG. 2A**, a pacemaker of the present invention shortens the AV/PV interval only after the occurrence of a prescribed number, e.g., three, consecutive R-waves 56a, 56b and 56c, each

R-wave evidencing a similar conduction time interval t_{AR} . Thus, as seen in **FIG. 2B**, the single occurrence of an R-wave 56a, or as shown in **FIG. 2C**, the occurrence of a pair of R-waves 56a and 56b, fails to cause the pacemaker to shorten the AV/PV interval.

Another method for reducing the tainting effects of ectopic R-waves relative to determining the natural

conduction time involves analyzing the morphology of an R-wave and, based on the morphology analysis, determining whether the R-wave is an ectopic R-wave before reducing the AV/PV interval. As seen in **FIG.**

5 **3A**, the morphology of a naturally occurring

R-wave 58 is measured and stored. Such stored R-wave is then compared against the morphology of other R-waves that occur. For example, a comparison between hypothetical R-waves, shown in **FIG. 3B**, and the

10 reference R-wave 58, shown in **FIG. 3A**, would result in a match only for the first measured R-wave 59.

One application of the invention, and the principal application described hereinafter in more detail, is to treat patients suffering from

15 hypertrophic obstructive cardiomyopathy (HOCM). In such instances, the AV (or PV) interval of the pacemaker is first set to a value that is less than the natural conduction time, t_{AR} , by a prescribed amount Δ , thereby assuring that ventricular pacing (in
20 the absence of a decreasing t_{AR}) will preempt a natural depolarization (R-wave). Upon the detection of non-ectopic R-waves, indicating a decreasing natural conduction time t_{AR} , the AV/PV interval is decreased to assure ventricular pacing. Such ventricular pacing
25 advantageously alters the mechanical contraction sequence of the cardiac tissue so as to favorably improve the performance of the heart.

In order to explain the effect of a cardiomyopathy, reference is made to **FIGS. 4 and 5**,

30 where there is shown a schematic cross-sectional representation of a healthy heart 12 (**FIG. 4**) and a heart 12' suffering from a HOCM (**FIG. 5**). Referring first to **FIG. 4**, the healthy heart 12 includes four chambers, a right atrium 302, a right ventricle 304, a
35 left atrium 306, and a left ventricle 308. The right

side of the heart is separated from the left side of the heart by a wall of tissue, or septum 300 and 324. The AV node 310 is located in the intra-atrial septum 300. A bundle of His 316 also lies in the intra-atrial septum 300, and provides a path through which a suitable stimulus is carried from the AV node 316 to the ventricular tissue near the apex of the heart. In response to such a stimulus, the ventricular tissue begins to contract, commencing along the intra-ventricular septum 324 in the direction of the arrows 326. Such contraction triggers a particular type of mechanical or physical action within the heart that forces the blood held in the ventricles out appropriate arteries (not shown) to desired body tissue locations.

In Fig. 5, there is shown a schematic representation of a heart 12' that suffers from hypertrophic obstructive cardiomyopathy. The left ventricular muscle mass 322, is even thicker than the muscle of the normal heart with the intra-ventricular septum 324' being the thickest of all causing it to bulge into the left ventricular cavity. With normal electrical conduction, the intra-ventricular septum 324' further thickens and acts like a splint with the rest of the ventricle bending around this structure. Effectively, this causes a relative or dynamic obstruction to the ejection of blood into the aorta. By artificially inducing a disordered sequence of activation, the vigor of contraction is reduced decreasing the degree of obstruction and improving cardiac function.

The improvement in the cardiac output is caused, at least in part, by the fact that the stimulated contraction of the heart 12' starts at a different location (i.e., at or near the electrode tip 17 of a ventricular pacing lead 16), than does a natural

apical contraction. The resulting stimulated contraction is represented in FIG. 5 by the arrow 330. Such stimulated contraction advantageously produces a different mechanical or physical sequence of the ventricular contraction than is achieved through a natural apical contraction 44 (FIG. 4). Such different mechanical sequence, in turn, creates pumping forces within the contracting muscle tissue that enhances the cardiac output. Further, for some patients, a continued treatment using preemptive ventricular pacing as described herein results in a marked improvement in cardiac output over time. That is, the continued application of a ventricular stimulus so as to cause a different mechanical contraction sequence appears to cause a thinning of the thick cardiomyopic walls of the heart, and a strengthening of the ventricular muscle tissue, thereby effecting not only a treatment for a cardiomyopathy, but also a cure.

To this end, the present invention determines the natural conduction time between a P-wave (evidencing depolarization of the atria) and a subsequent R-wave, or P-R interval, and sets the PV interval of the pacemaker to be a prescribed amount less than such P-R interval. Alternatively, should the atria of the patient also require stimulation, the invention determines the paced conduction time between an atrial stimulation pulse, (A-pulse) and a subsequent R-wave, or A-R interval, and sets the AV interval of the pacemaker to be a prescribed amount less than such A-R interval. In this manner, the pacemaker always delivers a V-pulse at the conclusion of the PV or AV intervals, which (for treating HOCM) is less than the natural conduction time (P-R or A-R interval), and hence before the ventricles attempt to contract on their own. In determining the A-R interval, any of

the techniques described above for distinguishing between an ectopic R-wave and a natural non-paced R-wave may be used.

Advantageously, the present invention may be implemented using a wide variety of dual-chamber pacemaker configurations and pacemaker hardware. Any pacemaker configuration that allows the pacemaker to determine the natural conduction time interval, i.e., the A-R interval, untainted by ectopic R-waves, and which further allows the AV or PV intervals to be automatically set to a value that is a prescribed amount less than the A-R or P-R conduction-time intervals, may be used to implement the invention. The description that follows is only exemplary of one such configuration.

A Representative Microprocessor-Based Pacemaker

To better understand how the present invention may be practiced, it will be helpful to review the main components, and basic operation, of a pacing system. Accordingly, the following overview of a pacemaker is presented.

In FIG. 6, a functional block diagram of a dual-chamber pacemaker 10 of a type with which the present invention may be used is shown. The pacemaker 10 is coupled to a heart 12 by way of leads 74 and 76, the lead 74 having an electrode 75 that is in contact with one of the atria of the heart, and the lead 76 having an electrode 77 that is in contact with one of the ventricles of the heart. The leads 74 and 76 are electrically and physically connected to the pacemaker 10 through a connector 73 that forms an integral part of the housing wherein the circuits of the pacemaker are housed.

The connector 73 is electrically connected to a protection network 78, which network 78 electrically

protects the circuits within the pacemaker 10 from excessive shocks or voltages that could appear on the electrodes 75 and/or 77 in the event such electrodes were to come in contact with a high voltage signal, e.g., from a defibrillator shock.

The leads 74 and 76 carry stimulating pulses to the electrodes 75 and 77 from an atrial pulse generator (A-PG) 79 and a ventricular pulse generator (V-PG) 80, respectively. Further, electrical signals from the atria are carried from the electrode 75, through the lead 74, to the input terminal of an atrial channel sense amplifier (P-AMP) 82; and electrical signals from the ventricles are carried from the electrode 77, through the lead 76, to the input terminal of a ventricular channel sense amplifier (R-AMP) 84. Similarly, electrical signals from both the atria and ventricles are applied to the inputs of an IEGM (intracardiac electrogram) amplifier 85. The amplifier 85 is typically configured to detect an evoked response from the heart 12 in response to an applied stimulus, thereby aiding in the detection of capture. (Capture occurs when an electrical stimulus applied to the heart is of sufficient energy to depolarize the cardiac tissue, thereby causing the heart muscle to contract, or in other words, causing the heart to beat. Capture does not occur when an electrical stimulus applied to the heart is of insufficient energy to depolarize the cardiac tissue.)

The dual-chamber pacemaker 10 is controlled by a control system 86 that typically includes a microprocessor programmed to carry out control and timing functions. The control system 86 receives the output signals from the atrial (P-AMP) amplifier 82 over signal line 88. Similarly, the control system 86 receives the output signals from the ventricular (R-AMP) amplifier 84 over signal line 90, and the

output signals from the IEGM amplifier 85 over signal line 91. These output signals are generated each time that a P-wave, an R-wave, or an evoked response is sensed within the heart 28. The control system 86
5 also generates trigger signals that are sent to the atrial pulse generator (A-PG) 79 and the ventricular pulse generator (V-PG) 80 over signal lines 92 and 94, respectively. These trigger signals are generated each time that a stimulation pulse is to be generated
10 by the respective pulse generator 79 or 80. The atrial trigger signal is referred to simply as the "A-trigger", and the ventricular trigger signal is referred to as the "V-trigger". During the time that either an A-pulse or V-pulse is being delivered to the
15 heart, the corresponding amplifier, P-AMP 82 and/or R-AMP 84, is typically disabled by way of a blanking signal presented to these amplifiers from the control system over signal lines 96 and 98, respectively. This blanking action prevents the amplifiers 82 and 84
20 from becoming saturated from the relatively large stimulation pulses that are present at their input terminals during this time. This blanking action also helps prevent residual electrical signals present in the muscle tissue as a result of the pacer stimulation
25 from being interpreted as P-waves or R-waves.

Still referring to FIG. 6, the pacer 10 also includes a memory circuit 100 coupled to the control system 86 over a suitable data/address bus 102. This memory circuit 100 allows certain control parameters,
30 used by the control system 86 in controlling the operation of the pacemaker, to be programmably stored and modified, as required, in order to customize the pacer's operation to suit the needs of a particular patient. Further, data sensed during the operation of
35 the pacer may be stored in the memory 100 for later retrieval and analysis.

The memory 100 of the pacemaker 10 may take many forms, and may be subdivided into as many different memory blocks or sections (addresses) as needed in order to allow desired data and control information to be stored. A feature of the present invention is the ability to store a relatively large amount of sensed data as a data record, which data record may then be used to guide the operation of the device, e.g., whether a given R-wave is an ectopic R-wave or a naturally occurring R-wave. Other parameters, of course, in addition to (or in lieu of) R-wave morphology data or conduction time data, may be similarly sensed, stored, averaged (or otherwise processed), and then used for comparison purposes. Advantageously, modern memory devices allow for the storage of large amounts of data in this manner.

A clock circuit 103 directs an appropriate clock signal(s) to the control system 86, as well as to any other needed circuits throughout the pacemaker 10 (e.g., to the memory 100) by way of clock bus 105.

A telemetry/communications circuit 104 is further included in the pacer 10. This telemetry circuit 104 is connected to the control system 86 by way of a suitable command/data bus 106. In turn, the telemetry circuit 104, which is included within the implantable pacer 10, may be selectively coupled to an external programming device 108 by means of an appropriate communication link 110, which communication link 110 may be any suitable electromagnetic link, such as an RF (radio frequency) channel, a magnetic link, an inductive link, an optical link, and the like. Advantageously, through the external programmer 108 and the communication link 110, desired commands may be sent to the control system 86. Similarly, through this communication link 110 and the programmer 108, data (either held within the control system 86, as in a data latch or stored within the memory 100) may be

remotely received from the pacer 10. In this manner, noninvasive communications can be established with the implanted pacer 10 from a remote, non-implanted, location.

5 The pacer 10 additionally includes a battery 93, which provides operating power to all of the circuits of the pacer 10 via a POWER signal line 95. The voltage of the pacemaker battery 93 on the POWER
10 signal line 95 is monitored by appropriate monitoring circuits within the control system 86, or elsewhere within the pacemaker 10, in order to ascertain whether the battery voltage has dropped below a predefined recommended replacement time (RRT) threshold.

 It is noted that the pacer 10 in **FIG. 6** is
15 referred to as a dual-chamber pacemaker because it interfaces with both the atria and the ventricles of the heart. Those portions of the pacer 10 that interface with the atria, e.g., the lead 74, the P-wave sense amplifier 82, the A-PG 78, and
20 corresponding portions of the control system 86, are commonly referred to as the "atrial channel". Similarly, those portions of the pacer 10 that interface with the ventricles, e.g., the lead 76, the R-wave sense amplifier 84, the V-pulse generator 80,
25 and corresponding portions of the control system 86, are commonly referred to as the "ventricular channel".

 As needed for rate-responsive applications, the pacemaker 10 further includes at least one sensor 112 that is connected to the control system 86 of the
30 pacemaker 10 over a suitable connection line 114. While this sensor 112 is illustrated in **FIG. 6** as being included within the pacer 10, it is to be understood that the sensor may also be external to the pacer 10, yet still be implanted within or carried by the
35 patient. Two common types of sensors include an activity sensor, such as a piezoelectric crystal, that

is mounted to the case of the pacemaker and an accelerometer mounted on the circuit substrate. Other types of sensors are also known, such as sensors that sense the oxygen content of blood, respiration rate, pH of blood, body motion, and the like. The type of sensor used is not critical to the present invention.

Any sensor capable of sensing a parameter relatable to the rate at which the heart should be beating can be used. Such sensors are used with "rate-responsive" pacemakers in order to adjust the basic rate (pacing cycle) of the pacer in a manner that tracks the physiological needs of the patient. To this end, the control system 86 (when operating in a rate-responsive mode) receives output signals from the sensor 112 and converts them to a sensor-indicated-rate (SIR) signal which is used by the control system 86 to set the pacing interval of the pacemaker.

The pacemaker 10 further includes magnet detection circuitry 87, coupled to the control system 86 over signal line 89. It is the purpose of the magnet detection circuitry 87 to detect when a magnet is placed over the pacemaker, which magnet may be used by a physician or other medical personnel to perform various reset functions of the pacemaker 10, and/or to signal the control system 86 that an external programmer 108 is in place to receive data from, or send data to, the pacemaker memory 100 or control system 86 through the telemetry communications circuits 104.

The telemetry or communications circuit 104 may be of conventional design, such as is described in U.S. Patent No. 4,944,299, or as is otherwise known in the art. Similarly, the external programmer 108 may be of any suitable design known in the art, such as is described in U.S. Patent No. 4,809,697. Likewise, the memory circuit 100, and the circuits utilized in the

atrial and ventricular channels may all be of common design as is known in the pacing art.

The control system 86 may be realized using a variety of different techniques and/or circuits. The
5 preferred type of control system 86 is a microprocessor-based control system, as described, e.g., in FIGS. 6-8 and accompanying text of U.S. Patent No. 5,476,487, incorporated herein by reference. It is noted, however, that the control
10 system 86 could also be realized using a state machine. Indeed, any type of control circuit or system could be employed for the control system 86. Those of skill in the implantable medical device art, given the teachings presented herein, should thus be able to
15 fashion numerous different types of control systems or circuits that achieve the desired stimulus pulses.

A representative alternative type of control system that could be used with the invention, for example, is the microprocessor-based control system
20 described in U.S. Patent No. 4,940,052, entitled "Microprocessor Controlled Rate-Responsive Pacemaker Having Automatic Rate Response Threshold Adjustment". Reference is also made to U.S. Patent Nos. 4,712,555 and 4,944,298, wherein a state-machine type of
25 operation for a pacemaker is described; and U.S. Patent No. 4,788,980, wherein the various timing intervals used within the pacemaker and their inter-relationship are more thoroughly described. The '052, '555, '298 and '980 patents are all incorporated
30 herein by reference.

Additional details relative to a particular multi-mode, microprocessor-based, pacemaker that may be utilized in order to implement the present invention are found in the previously-referenced U.S.
35 Patent No. 5,476,487. As indicated above, however, the present invention is not limited by a particular

pacemaker design, but may be implemented using any pacemaker design wherein operating parameters of the pacemaker and/or the patient/pacemaker interface may be monitored, measured, and stored, and wherein the
5 timing of ventricular stimuli may be implemented in the manner described herein.

Of primary significance to the present invention is the manner in which the AV interval (or PV interval) is adaptively adjusted to a value related to
10 the measured natural conduction time of the patient. The manner in which this is done is illustrated in the flowcharts of FIGS. 7 and 8. FIG. 7 illustrates the operation of the pacemaker in absence of an R-wave verified to be produced by natural AV conduction; that
15 is, a series of a predetermined or programmable number of consecutive R-waves (e.g., preferably three or more) or at least one R-wave of appropriate morphology. FIG. 8 illustrates the operation of the pacemaker in the presence of R-waves produced by
20 natural AV conduction.

As seen in FIG. 7, the method starts by setting the initial values needed by the pacemaker to carry out, e.g., DDD or DDDR pacing (block 502). (DDD and other pacing modes are described in U.S. Patent No.
25 5,237,992, which patent is incorporated herein by reference.) Such values are, for the most part, no different than those used when performing conventional DDD or DDDR pacing, and include such values as an initial pacing rate (from which an appropriate atrial
30 escape interval is determined), an initial value for the AV interval, blanking period values, maximum pacing rate values, stimulation pulse amplitudes and widths, and the like. In accordance with the present invention, such initial values also include a minimum
35 and maximum value for the AV (or PV) interval, plus a prescribed time difference between the natural

conduction time of the patient and the pacemaker-defined AV (or PV) interval. In some embodiments of the invention, it may also be important to specify the difference between the A-R interval and a P-R
5 interval, where the A-R interval is the natural conduction time as measured from the delivery of an A-pulse to the subsequent occurrence of an R-wave, and the P-R interval is the natural conduction time as measured from the occurrence of a P-wave to the
10 subsequent occurrence of an R-wave.

Once the initial values needed to carry out DDD or DDDR pacing have been set, the specified DDD or DDDR pacing is carried out (block 504) in conventional manner, one cardiac cycle at a time, using the
15 programmed values. At some point in a cardiac cycle associated with such DDD or DDDR pacing, an R-wave resulting from natural AV conduction will occur; an ectopic R-wave may occur; or a number of consecutive cardiac cycles will go by without the occurrence of an
20 R-wave.

The occurrence of R-waves may indicate the depolarization of the ventricles as a result of a natural or native conduction time shorter than the presently existing AV (or PV) interval of the
25 pacemaker, or the R-wave may be a single ectopic R-wave. When preemptive ventricular pacing is desired, (i.e., when treating HOCM), R-waves resulting from natural AV conduction may indicate that the pacemaker-defined AV (or PV) interval needs to be decreased.

30 To prevent an isolated R-wave or ectopic beat from prematurely shortening the AV interval, the control system will not determine a new natural conduction time (block 506) until a specified criteria is met, preferably, a predetermined (or programmable)
35 number of consecutive cardiac cycles must occur without an R-wave, thereby indicating a true lengthening of the natural conduction time.

Alternately, other criteria could simply include the passage of a prescribed number of cardiac cycles (e.g., between 8 and 2048), or the mere passage of time (e.g., once every 10-15 minutes) may be used to
5 signal a need to determine the natural conduction time of the patient (block 506). In another embodiment, the R-wave must have a predetermined (or pre-characterized) morphology to distinguish true conducted R-waves from ectopic R-waves.

10 Accordingly, as soon as an appropriate criteria is met, then the natural conduction time, t_{AR} , is determined or remeasured (block 506).

The natural conduction time is determined as the time interval between the most recent atrial activity,
15 which would be either a P-wave or an A-pulse, and an R-wave. That is, the native or natural (note, as used herein, "native" and "natural" are used as synonyms) begins with the occurrence of atrial activity, and ends with the occurrence of an R-wave. If the most
20 recent atrial activity was a P-wave, then the conduction time measured is t_{PR} . If the most recent atrial activity was an A-pulse, then the conduction time measured is t_{AR} . (Note, heretofore t_{AR} has been used to refer generically to the natural conduction
25 time, whether technically t_{PR} or t_{AR}).

In order to assure that the determination of t_{AR} or t_{PR} accurately reflects the true natural conduction time (and not tainted by an ectopic R-wave), the natural conduction time over several cardiac cycles
30 may be measured, and then an average, mean, or other appropriate combination, of t_{AR} or t_{PR} over the prescribed number of cycles can be computed.

If an R-wave fails to occur for a prescribed number of cardiac cycles, then that provides an
35 indication that perhaps the natural conduction time has increased, and that there is a need to increase

the AV (or PV) interval so that it is not too different than the natural conduction time.

Once such need is identified, an appropriate adjustment to the AV (or PV) interval of the pacemaker can be made, as required (block 508). That is, once a determination is made that the natural conduction time has either decreased or increased (block 506), the AV (or PV) interval of the pacemaker is then set to a value that is just less than the determined natural conduction time. This is done by setting the AV (or PV) interval to be equal to the most recent t_{AR} (or t_{PR}) determination minus Δ , where Δ is a programmable value, typically ranging from 5-100 ms. For example, when the pacemaker is used to treat HOCM, and when it appears that the natural conduction time has decreased (as is most often the case when an R-wave has been sensed), then the AV (or PV) interval is set to t_{AR} (or t_{PR}) minus Δ .

After the AV (or PV) intervals have been set to be less than the determined conduction time t_{AR} (or t_{PR}) at block 508, then a determination is made as to whether DDD or DDDR pacing is to continue (block 510). If not, then the method terminates (block 512). If so, then the method continues (block 504) by performing the DDD or DDDR pacing for the next cardiac cycle using the adjusted values of the AV (or PV) interval.

In FIG. 8, a more detailed flowchart is illustrated that shows one technique for determining or measuring the natural conduction time of the patient (block 506 in FIG. 7), and adjusting the AV (or PV) intervals accordingly (block 508 in FIG. 7). It is noted that the method shown in FIG. 7, as well as the method shown in FIG. 8, are directed primarily to using the pacemaker to treat HOCM (i.e., are

directed to a method that provides preemptive ventricular pacing). However, such methods may be readily adapted by those of skill in the art for other applications of the invention.

5 In FIG. 8, the programmed values needed to carry out DDD or DDDR pacing are programmed into the pacemaker in conventional manner (block 520). In accordance with the present invention, such programmed values include the number of cardiac cycles that must
10 occur without an R-wave before the AV (or PV) interval is increased, the amount of such increase, an initial value for the natural conduction time t_{AR} (or t_{PR}), or an indication of a technique for determining such initial values, the difference X and/or Y between the
15 natural conduction times and the AV (or PV) intervals, and the like (block 522). Once the initial values of t_{AR} or (t_{PR}) have been determined, then the value of the AV (or PV) interval is set to be a specified amount (e.g., Δ) less than t_{AR} or t_{PR} (block 524).

20 With the AV (or PV) interval set to an initial value, the DDD or DDDR pacing cycle commences using such value, plus the other programmed values (block 526). If an R-wave is sensed during the pacing cycle (block 528), then that may signal that the natural
25 conduction time t_{AR} (or t_{PR}) is shorter than the pacemaker-defined AV (or PV) interval. As discussed in more detail below, the occurrence of an R-wave is verified by a method of the present invention, i.e., the occurrence a number of consecutive R-waves (e.g. a
30 programmable number, preferably greater than three) or, alternatively, the occurrence of an R-wave having the appropriate morphology.

The occurrence of a verified R-wave indicates the end of the conduction time t_{AR} (or t_{PR}), and thus
35 permits a measurement of t_{AR} (or t_{PR}) to be completed (block 530), preferably based on a average of a at

least three or a programmable number of naturally
conducted R-waves. The measured value of t_{AR} (or t_{PR}) is
then used as a basis for decreasing the AV (or PV)
interval (block 532). The AV interval is set to $t_{AR}-X$,
5 where X is a parameter having a programmable value, a
fixed value, or an adaptive value based on a
percentage of the heart rate. Similarly, the PV
interval is set to $t_{PR}-Y$, where Y is a parameter having
a programmable value, a fixed value, or an adaptive
10 value based on a percentage of the heart rate.

Alternatively, the AV (or PV) interval can be
decreased by a programming interval Δ below the
existing AV (or PV) interval. Accordingly, the AV (or
PV) interval is decreased by the programmable interval
15 Δ without immediate measurement of the natural
conduction time t_{AR} or t_{PR} (block 532').

In some embodiments of the invention, t_{PR} and t_{AR}
are measured separately, and separate values are
programmed or otherwise determined for the parameters
20 X and Y. Thus, in such embodiments, t_{PR} and the
resulting PV interval, and t_{AR} and the resulting AV
interval, are totally independent of each other. In
other embodiments, one of t_{AR} or t_{PR} is determined,
e.g., whichever happens to occur first, and the other
25 is computed as a function of the measured value. In
such embodiments, t_{AR} is set to be a prescribed number
of ms greater than t_{PR} . In such embodiments, there is
thus a prescribed relationship between t_{AR} and t_{PR} and
the resulting AV and PV intervals. For most purposes
30 relating to the description of the present invention,
one of the AV (or PV) intervals, or one of the
conduction times t_{AR} (or t_{PR}), is all that is expressly
referenced, and it is assumed that the other can be
determined in an appropriate manner. Techniques for
35 measuring the natural conduction time of a patient are

described in U.S. Patent No. 5,340,361, already incorporated herein by reference.

In another embodiment, the occurrence of ectopic R-waves that fall within a limited time window of, e.g., 50 ms, is highly unlikely. Thus, if an average conduction time t_{AR} or (t_{PR}) for three consecutive R-waves is calculated, and the conduction time t_{AR} or (t_{PR}) for each R-wave falls within plus or minus 25 ms of the calculated average conduction time, then the occurrence of a R-wave resulting from natural A-V conduction is considered to have been verified (block 506). Other similar techniques relating to the occurrence and timing of a series of R-wave likewise may be used to verify the occurrence of R-waves resulting from natural A-V conduction.

The assessment of the morphology of an R-wave (or QRS complex) requires the pacemaker to learn the morphology of an R-wave resulting from a native or natural ventricular depolarization. An R-wave's morphology is sampled by the sense and IEGM amplifiers 428 (FIG. 9) which samples the R-wave at a fixed sample rate of, e.g., 256 bytes per second. Thus, in the learn mode, a native R-wave is sampled and the resulting data bytes stored. In an operational mode, a measured R-wave is sampled and converted to data bytes. The data from the measured R-wave is compared with the stored data from the native R-wave and a correlation figure calculated. If the correlation figure exceeds a predetermined threshold, then the measured R-wave is considered a verified R-wave, and if appropriate, the pacemaker performs the necessary AV/PV interval adjustments. If the correlation figure fails to exceed the predetermined threshold, the measured R-wave is considered an ectopic R-wave and ignored.

In still another embodiment, an R-wave may be verified (block 528) by analyzing the timing or the

morphology of the R-wave. To allow for changes in R-wave morphology over time, a physician also may have R-wave signals digitized and transmitted via a telemetry interface 110 (FIG. 6) to an external
5 programmer 108 where the signals are converted back to their original analog form for display. If desired, the physician may "reset" the stored native morphology. More specifically, the physician, using external program control of the microprocessor control
10 and timing circuit 86, temporarily increases the AV/PV interval to enable the occurrence of intact AV nodal conduction. A newly acquired R-wave is then transmitted to a display for the physician's approval. Upon approval, the new native morphology is stored as
15 the reference R-wave morphology. It is further possible to have multiple morphologies stored in memory, with a particular morphology being selected by the physician or automatically being selected based on the patient's physiological condition. Another
20 representative morphology system for use in implementing the pacemaker of the present invention is described in U.S. Patent No. 5,513,644, incorporated herein by reference.

Thus, it is seen that the present invention
25 provides an implantable pacemaker, and method of operating such a pacemaker, that can set the AV (or PV) interval of the pacemaker to a value that is derived from the measured natural conduction time of the patient. For treating HOCM, for example, such
30 pacemaker and method stimulates the cardiac tissue at a time in the cardiac cycle that is just prior to when natural depolarization of the cardiac tissue would otherwise cause a cardiac contraction, and in so
doing, creates a mechanical contraction sequence that,
35 for most patients, improves the cardiac output..

As further described above, it is seen that the invention provides a dual-chamber pacemaker, and method of operating such a dual-chamber pacemaker, that automatically sets and adjusts the pacemaker-defined AV (or PV) interval to a value that is generally just less than the natural conduction time of the patient. Such action advantageously causes a V-pulse to be generated and delivered to the ventricular muscle tissue at a time in the cardiac cycle that suits a particular purpose (e.g., treating a HOCM or providing demand pacing), while still maintaining the approximate cardiac timing set by the natural conduction time.

As also described above, and for use with HOCM treatment, it is seen that the invention provides a dual-chamber pacemaker, and method of operating such a pacemaker, that decreases the pacemaker-defined AV interval in response to sensing a verified R-wave (which verified R-wave evidences a shortened natural conduction time), and that automatically increases the pacemaker-defined AV interval in prescribed increments in response to not sensing an R-wave for a prescribed number of consecutive cardiac cycles (which failure to sense any R-waves may evidence a lengthening of the natural conduction time). Thus, advantageously, the pacemaker-defined AV interval is most always set to a value that is just somewhat less than the natural conduction time, regardless of whether the natural conduction time is increasing or decreasing, and the cardiac output of the patient is maximized.

While the invention herein disclosed has been described by means of specific embodiments and applications thereof, numerous modifications and variations could be made thereto by those skilled in the art without departing from the scope of the invention set forth in the claims.

CLAIMSWhat is claimed is:

1. A dual-chamber pacemaker providing atrio-
5 ventricular pacing in order to increase cardiac output
in a patient suffering from hypertrophic obstructive
cardiomyopathy, the dual-chamber pacemaker including
atrial sensing means for sensing P-waves in the
atrium, ventricular sensing means for sensing R-waves
10 in the ventricle, and pulse generating means for
generating an atrial stimulation pulse (A-pulse) in
the atrium in the absence of a sensed P-wave, and a
ventricular stimulation pulse (V-Pulse) in the
ventricle in the absence of a sensed R-wave within an
15 AV/PV interval, the dual-chamber pacemaker comprising:
measuring means for measuring A-R/P-R intervals,
wherein the A-R interval represents the
interval between an A-pulse and the R-wave,
and the P-R interval represents the interval
20 between a sensed P-wave and the R-wave,
whichever event occurs first;
means for processing the A-R and P-R intervals so
as to minimize the effects of ectopic R-waves
and to produce as an output a conduction time
25 interval corresponding to a naturally-
conducted R-wave; and
control means for adjusting the AV/PV interval to
a value that is less than the conduction time
interval.

30

2. The dual-chamber pacemaker, as defined in
Claim 1, wherein:
the processing means includes means for detecting
a first prescribed number of consecutive

cardiac cycles with a sensed R-wave in the AV/PV interval, the presence of such sensed R-waves indicating a true decrease in the conduction time interval; and

5 the control means includes means for decreasing the AV-PV interval by a first prescribed interval less than the conduction time interval in response to the detection of a true decrease in the conduction time

10 interval.

3. The dual-chamber pacemaker, as defined in Claim 2, wherein the first prescribed number of consecutive cardiac cycles with a sensed R-wave in the

15 AV/PV interval comprises a programmable number of consecutive cardiac cycles.

4. The dual-chamber pacemaker, as defined in Claim 2, wherein the first prescribed number of

20 consecutive cardiac cycles with a sensed R-wave in the AV/PV interval comprises at least three consecutive cardiac cycles.

5. The dual-chamber pacemaker, as defined in

25 Claim 2, wherein the processing means further comprises:

means for averaging the conduction time interval so as to minimize the effects of ectopic R-waves.

30

6. The dual-chamber pacemaker, as defined in Claim 5, wherein the average conduction time interval is adjusted to a new value only if the R-waves occur in the AV/PV interval in excess of a predetermined

time range about a current average conduction time interval.

7. The dual-chamber pacemaker, as defined in
5 Claim 6, wherein the predetermined time range
comprises at least 25 ms plus and minus about the
current average conduction time interval.

8. The dual-chamber pacemaker, as defined in
10 Claim 2, wherein the first prescribed interval
comprises an interval which is in the range of 5-100
ms.

9. The dual-chamber pacemaker, as defined in
15 Claim 8, wherein the first prescribed interval
comprises an interval which is at least 20 ms less.

10. The dual-chamber pacemaker, as defined in
Claim 1, wherein:
20 the processing means includes means for detecting
a second prescribed number of consecutive
cardiac cycles without a sensed R-wave in the
AV/PV interval, the absence of such sensed R-
waves indicating a true increase in the
25 conduction time interval; and
the control means includes means for increasing
the AV-PV interval by a second prescribed
interval in response to the detection of a
true increase in the conduction time
30 interval.

11. The dual-chamber pacemaker, as defined in
Claim 10, wherein the prescribed number of consecutive
cardiac cycles without a sensed R-wave in the AV/PV

interval comprises at least three consecutive cardiac cycles.

12. The dual-chamber pacemaker, as defined in Claim 10, wherein the prescribed number of consecutive cardiac cycles without a sensed R-wave in the AV/PV interval comprises a programmable number of consecutive cardiac cycles.

13. The dual-chamber pacemaker, as defined in Claim 12, wherein the prescribed number of consecutive cardiac cycles without a sensed R-wave wave in the AV/PV interval comprises a programmable number between 4 and 512 cardiac cycles.

15

14. A dual-chamber programmable pacemaker for enhancing cardiac performance of a patient suffering from hypertrophic obstructive cardiomyopathy, comprising:

20 sensing means for sensing atrial and ventricular contractions;
 measuring means for measuring a conduction time interval between atrial contractions, either a paced or an intrinsic P-wave, and an
25 intrinsic R-wave corresponding to a ventricular contraction;
 timing means for defining an AV/PV interval as the time between the atrial contraction and a ventricular stimulation pulse which occurs in
30 the absence of an R-wave;
 processing means for decreasing an initial value of the AV/PV interval by a predetermined time interval in response to at least three consecutive sensed R-waves, indicative of
35 repetitive naturally-conducted ventricular

contractions, and for increasing the AV/PV interval to its initial value after a predetermined number of ventricular contractions.

5

15. A dual-chamber programmable pacemaker for enhancing cardiac performance of a patient suffering from hypertrophic obstructive cardiomyopathy, comprising:

- 10 sensing means for sensing P-waves and R-waves, associated with natural atrial and ventricular contractions, respectively; means for storing morphology signals for a selected reference R-wave;
- 15 means for verifying, based on a comparison of the reference R-wave, whether a current R-wave has the morphology of a naturally-conducted R-wave or is the result of an ectopic R-wave; means for determining a natural conduction time
- 20 interval between an atrial event, either a P-wave or a paced atrial event, and a verified naturally-conducted R-wave; processing means for adjusting an AV/PV interval between the atrial event and a paced
- 25 ventricular contraction, to a value that is a prescribed time interval less than the natural conduction time interval.

16. A dual-chamber programmable pacemaker, as
- 30 defined in Claim 15, wherein the processing means, after shortening the AV/PV interval, returns the AV/PV interval to its initial value after a predetermined number of paced ventricular contractions.

17. The dual-chamber programmable pacemaker, as defined in Claim 10, wherein the prescribed time interval is 5-100 ms.

5 18. The dual-chamber programmable pacemaker, as defined in Claim 17, wherein the predetermined number of paced ventricular contractions is between 4 and 512.

10 19. The dual-chamber programmable pacemaker, as defined in Claim 15, wherein the processing means decreases the AV/PV interval after at least three consecutive verified R-waves occur.

15 20. The dual-chamber programmable pacemaker, as defined in Claim 15, wherein the processing means selects the reference R-wave from a plurality of R-wave morphologies stored in the processor.

20 21. The dual-chamber programmable pacemaker, as defined in Claim 20, wherein the processor selects the reference R-wave under external control through a telemetry link.

25 22. The dual-chamber programmable pacemaker, as defined in Claim 20, further comprising:

 a physiological sensor that generates a signal representing the patient's physiological condition; and

30 wherein the processing means automatically selects the reference R-wave based on the patient's physiological condition.

23. A method of providing atrio-ventricular
35 pacing in order to enhancing cardiac performance in a

patient suffering from hypertrophic obstructive cardiomyopathy, comprising the steps of:

- measuring A-R/P-R intervals, wherein the A-R interval represents the interval between an A-pulse and the R-wave, and the P-R interval represents the interval between a sensed P-wave and the R-wave, whichever event occurs first;
- processing the A-R and P-R intervals so as to minimize the effects of ectopic R-waves and to produce as an output a conduction time interval corresponding to a naturally-conducted R-wave;
- defining an adjustable AV/PV interval as the interval beginning upon the detection of the atrial activity and ending with a ventricular stimulation pulse which occurs in the absence of an intrinsic R-wave; and
- adjusting the AV/PV interval to a value that is less than the processed conduction time interval by a prescribed amount.

24. The method of providing atrio-ventricular pacing, as defined in Claim 23, wherein the adjusting step further comprises the step of:

decreasing the AV/PV interval by a first prescribed interval after at least three consecutive sensed R-waves.

25. The method of providing atrio-ventricular pacing, as defined in Claim 13, further comprising the step of:

increasing the AV/PV interval by a second prescribed interval after a prescribed number of consecutive cardiac cycles without a sensed R-wave.

26. A method of operating a dual-chamber programmable pacemaker for enhancing cardiac performance of a patient suffering from hypertrophic obstructive cardiomyopathy, comprising the steps of:

5 sensing an R-wave;

 comparing the sensed R-wave with a reference R-wave to verify that the sensed R-wave is the result of a conducted ventricular contraction and is not the result of an ectopic ventricular contraction;

10 measuring the conduction time interval based on an atrial event and a verified R-wave; and

 setting an AV/PV interval between an atrial event and a paced ventricular contraction to a value that is a prescribed time interval less than the measured conduction time interval.

15

27. The method of operating a dual-chamber programmable pacemaker, as defined in Claim 26, further comprising the step of:

20 decreasing the value of the AV/PV interval by a first predetermined time interval in response to at least three consecutive verified R-waves indicative of natural ventricular contractions.

25

28. The method of operating a dual-chamber programmable pacemaker, as defined in Claim 26, further comprising the step of:

30 selecting the reference R-wave from a plurality of R-wave morphologies stored in the processor.

29. The method of operating a dual-chamber programmable pacemaker, as defined in Claim 28, further comprising the steps of:

- 5 generating physiological signal representing the patient's physiological condition; and
automatically selecting the reference R-wave based on the patient's physiological condition.

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FIG. 1
PRIOR ART

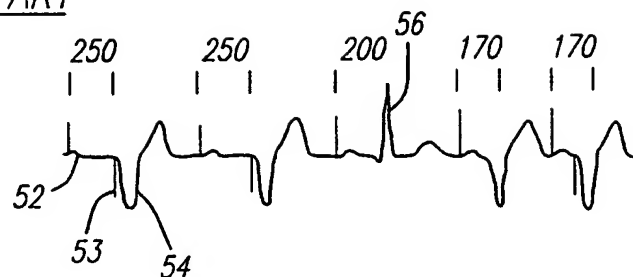


FIG. 2A

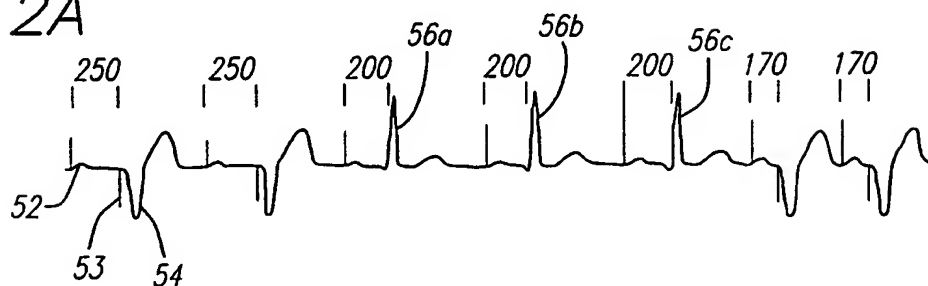


FIG. 2B

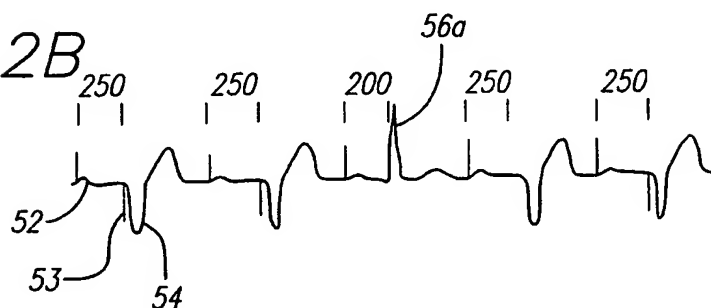
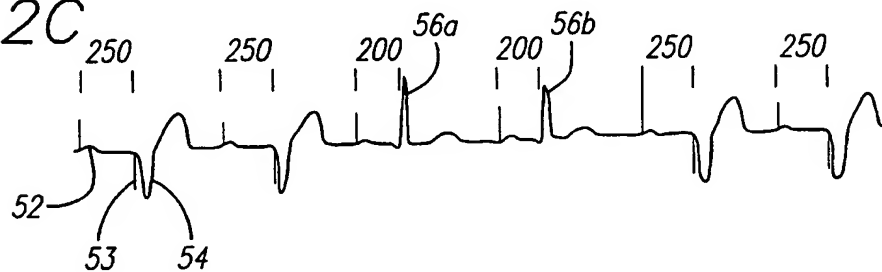


FIG. 2C



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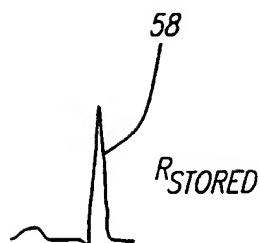


FIG. 3A

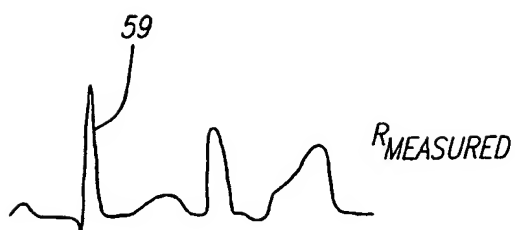
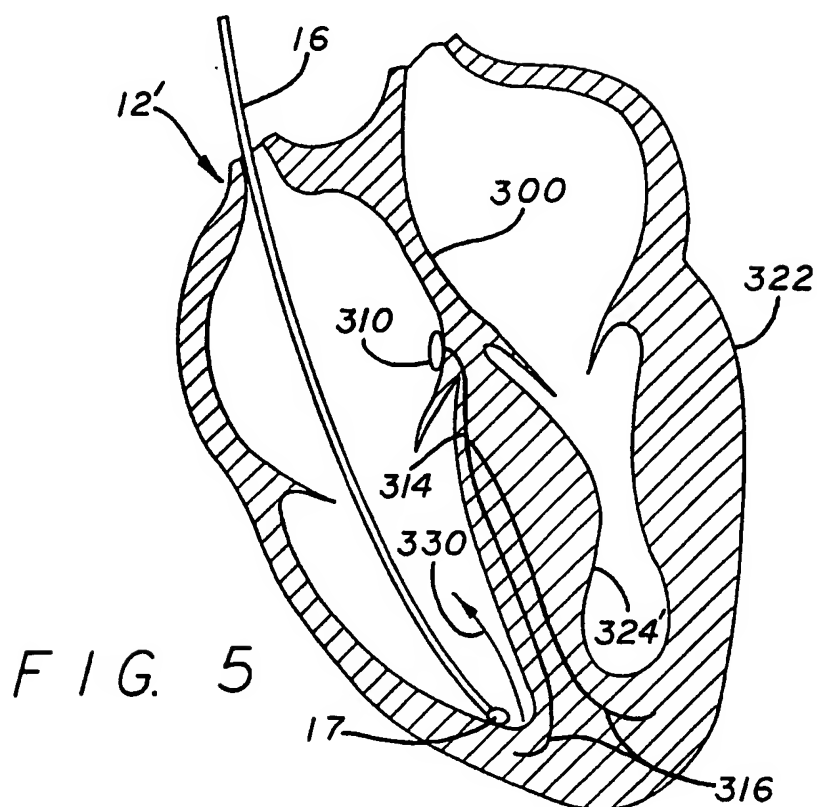
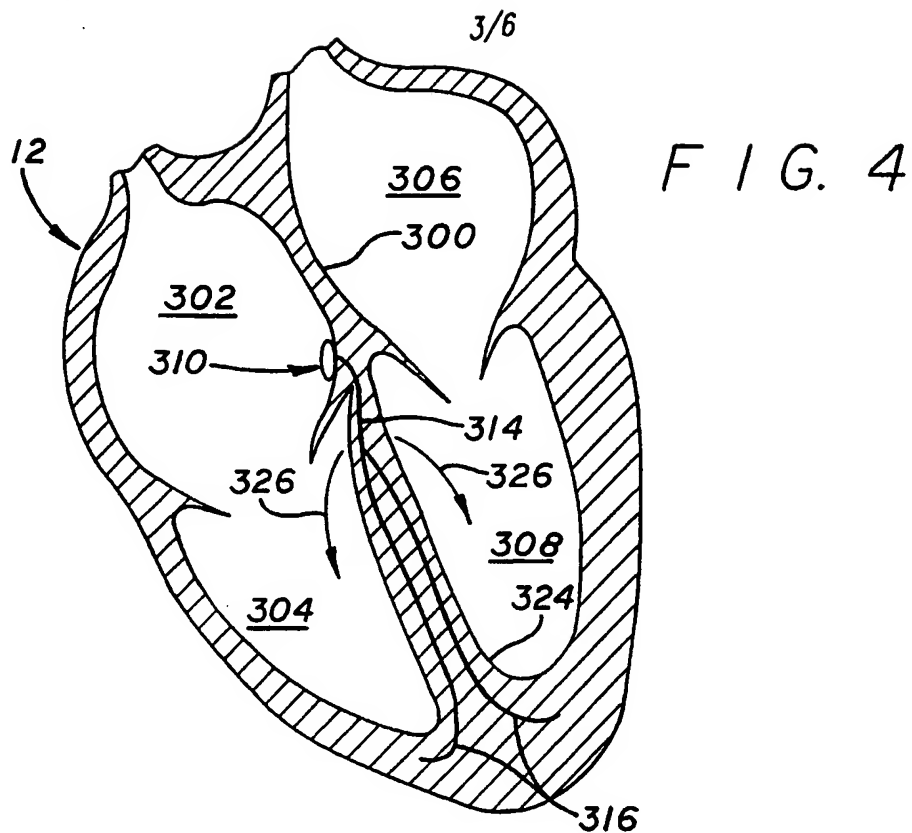


FIG. 3B



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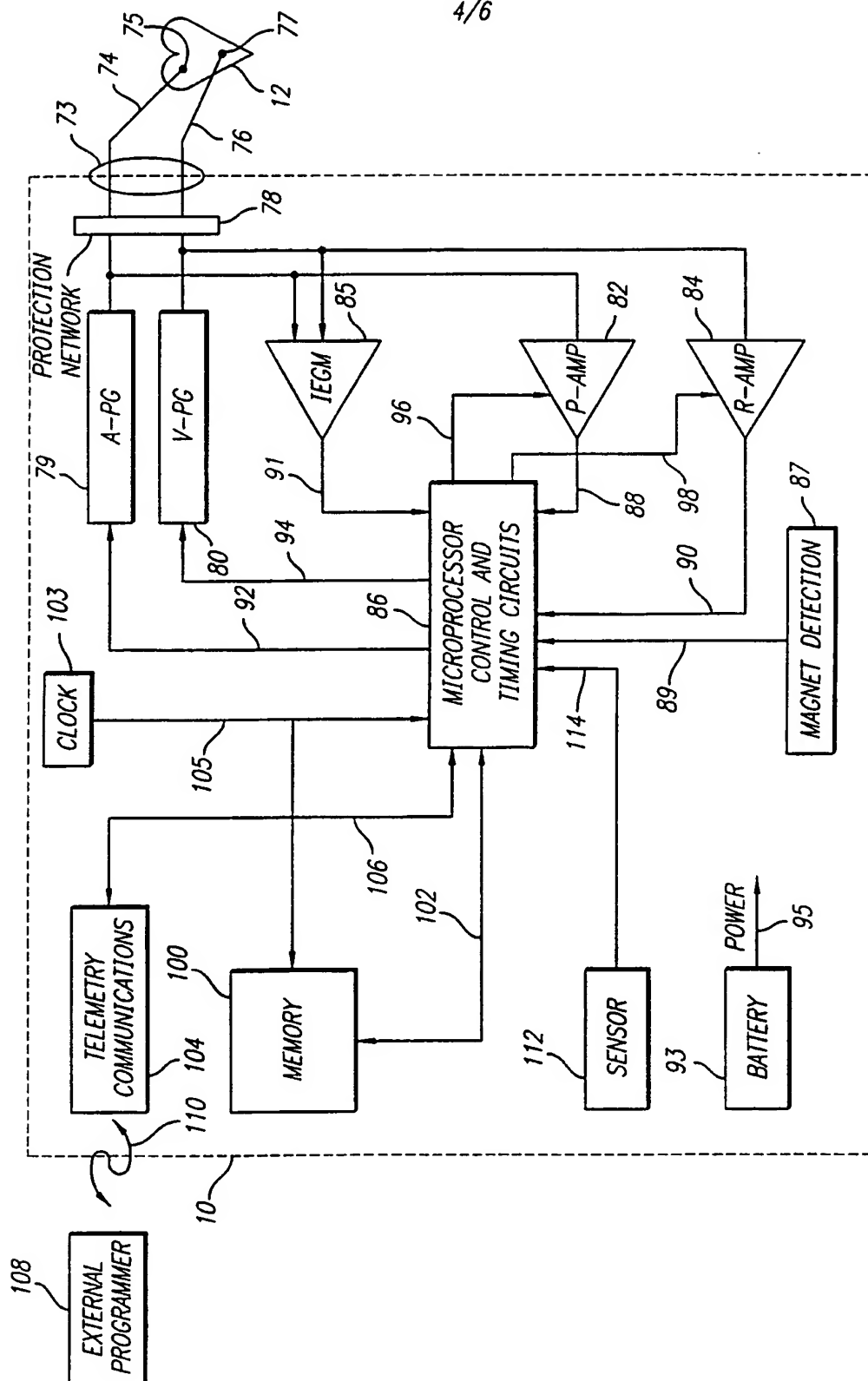
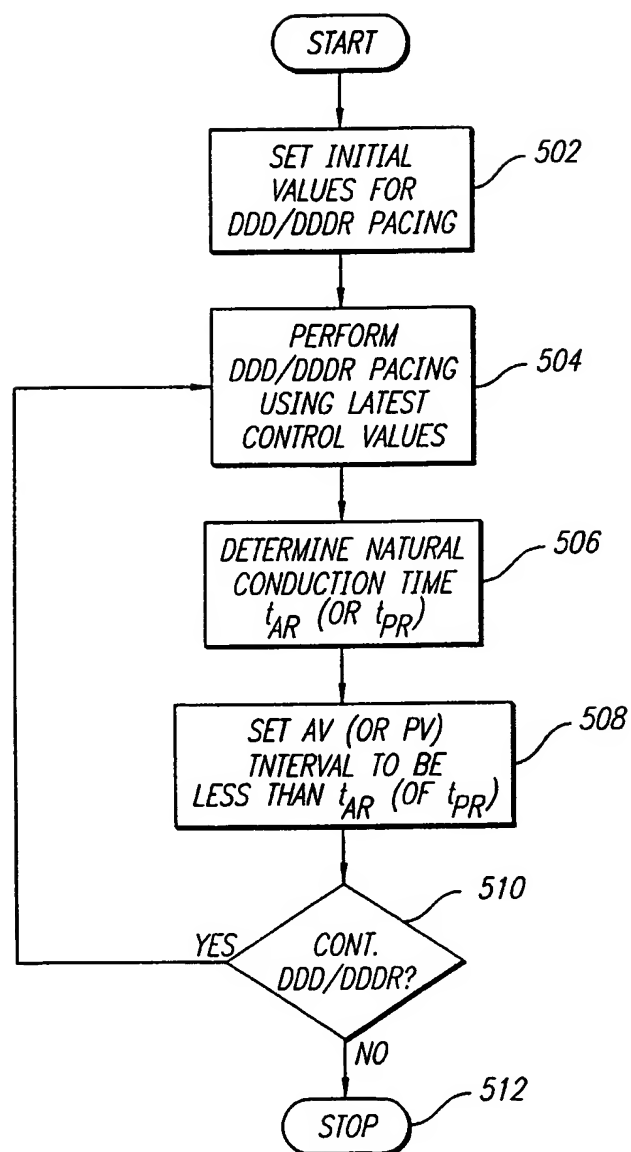


FIG. 6

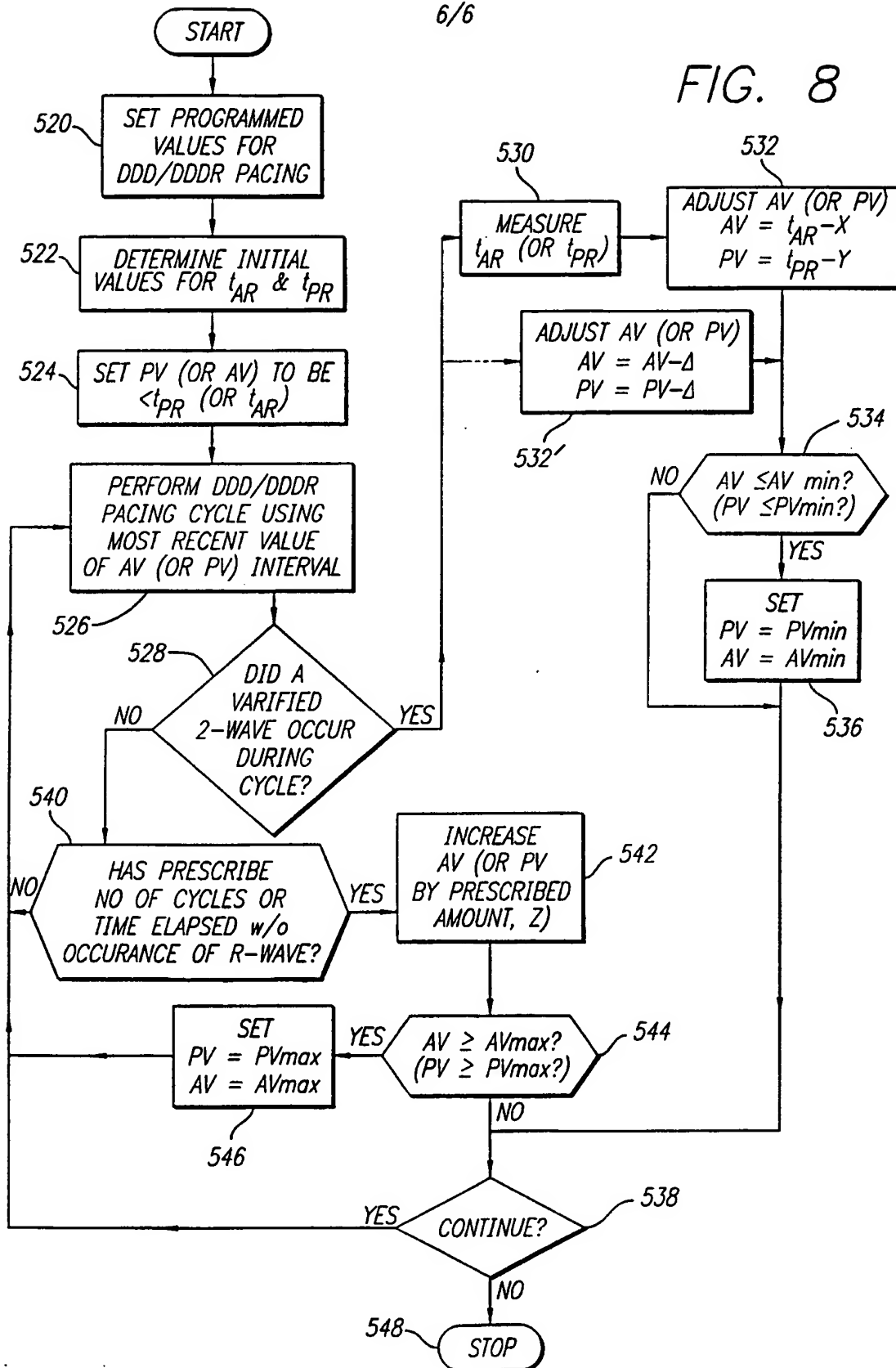
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FIG. 7



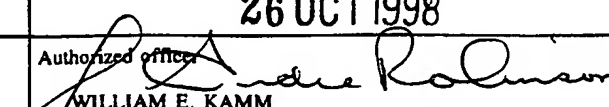
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FIG. 8



INTERNATIONAL SEARCH REPORT

International application No.
PCT/US98/17506

A. CLASSIFICATION OF SUBJECT MATTER		
IPC(6) : A61N 1/362 US CL : 607/009 According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED		
Minimum documentation searched (classification system followed by classification symbols) U.S. : 607/009, 013		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 5,527,347 A (SHELTON et al) 18 June 1996, entire document.	1-29
A	US 5,514,163 A (MARKOWITZ et al) 7 May 1996, entire document.	1-29
<input type="checkbox"/> Further documents are listed in the continuation of Box C. <input type="checkbox"/> See patent family annex.		
* Special categories of cited documents:	*T*	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
A document defining the general state of the art which is not considered to be of particular relevance	*X*	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
E earlier document published on or after the international filing date	*Y*	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*A*	document member of the same patent family
O document referring to an oral disclosure, use, exhibition or other means		
P document published prior to the international filing date but later than the priority date claimed		
Date of the actual completion of the international search 02 OCTOBER 1998	Date of mailing of the international search report 26 OCT 1998	
Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Facsimile No. (703) 305-3230	Authorized officer:  WILLIAM E. KAMM Telephone No. (703) 308-2994	